

EU Certificate

for the assessment of the
quality management system



according to Medical Device Regulation (EU) 2017/745 Annex IX Chapter I+III

As a Notified Body of the European Union DEKRA Certification GmbH certifies, that the manufacturer

VITA Zahnfabrik H. Rauter GmbH & Co. KG

Single Registration Number (SRN): DE-MF-000005906

Spitalgasse 3, 79713 Bad Säckingen, Germany

applies a quality management system according to Annex IX Chapter I+III of the Medical Device Regulation (EU) 2017/745 for the medical devices listed in the annex. This certificate is based on the assessments listed in CNo50069-01 and is only valid in conjunction with the successful completion of the annual surveillance audits.

EU Certificate no.: 50069-60-03-00

Certificate valid from:

2026-10-12

Certificate valid to:

2031-10-11

Previous certificate no. 50069-60-02-00, issued on 2025-05-20

Digitally signed by Markus RAINER
Kopf
Date: 2026-10-06 15:28:14+02:00



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de

BS-MDR-092

DEKRA Certification GmbH, Stuttgart
Notified Body ID number: 0124

Annex to the EU Certificate no. 50069-60-03-00

Following devices/device categories are included in this certificate:

Class IIa

- Artificial teeth for removable and fixed dentures
- Dental CAD/CAM Materials
 - Polymer, implantable and non-implantable
 - Ceramic, implantable
 - Polymer-Ceramic, implantable
- Dental Veneering Materials
 - Polymer, implantable
 - Ceramic, implantable

Change(s) to previous certificate: Reassessment